



The Patent Office
Concept House
Cardiff Road
Newport
South Wales
NP10 8QQ

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

I also certify that the attached copy of the request for grant of a Patent (Form 1/77) bears a correction, effected by this office, following a request by the applicant and agreed to by the Comptroller-General.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

Signed *Andrew Gentry*

Dated - 2 APR 2001

Request for grant of a patent
(See the notes on the back of this form. You can also get
an explanatory leaflet from the Patent Office to help
you fill in this form)



10 SEP 1998

The Patent Office
Cardiff Road
Newport
Gwent NP9 1RH

1. Your reference

AGS/PS/P21245GB

2. Patent application number
(The Patent Office will fill in this part)

Fil/77
CF. 16/2/99

3. Full name, address and postcode of the or of
each patent applicant (underline all surnames)

OXAGEN LIMITED
3 WORCESTER STREET
OXFORD OX1 2PZ

ERASMUS UNIVERSITEIT
ROTTERDAM
BURGEMEESTER OUDLAAN 50
3062 PA ROTTERDAM
THE NETHERLANDS

Patents ADP number (if you know it)

00434191001

If the applicant is a corporate body, give the
country/state of its incorporation

4. Title of the invention

METHOD

5. Name of your agent (if you have one)

KILBURN & STRODE
20 RED LION STREET
LONDON
WC1R 4PJ

Patents ADP number (if you know it)

125001

6. If you are declaring priority from one or more
earlier patent applications, give the country and
the date of filing of the or each of these earlier
applications and (if you know it) the or each
application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise
derived from an earlier UK application,
give the number and the filing date of
the earlier application

Number of earlier application number

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right
to grant of a patent required in support of
this request? (Answer 'Yes' if:

YES

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an
applicant, or
- c) any named applicant is a corporate body.
See note (d))

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form.
Do not count copies of the same document

Continuation sheets of this form

Description	:	17	/
Claim(s)	:	3	/
Abstract	:		/
Drawing(s)	:	5	x 5 /

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right
to grant of a patent (Patents Form 7/77)

Request for preliminary examination
and search (Patents Form 9/77)

Request for substantive examination
(Patents Form 10/77)

Any other documents
(please specify)

I/We request the grant of a patent on the basis of this application.

11.

Signature

Date

10 September 1998

Kilburn & Strode

Kilburn & Strode

12. Name and daytime telephone number of person to contact in the United Kingdom

Ms. Punita Shah
Tel: 0171-539 4200

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

a) If you need help to fill in this form or you have any questions, please contact the Patent Office on 0645 500505.

b) Write your answers in capital letters using black ink or you may type them.

c) If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.

d) If you have answered 'Yes' Patents Form 7/77 will need to be filed.

e) Once you have filled in the form you must remember to sign and date it.

f) For details of the fee and ways to pay please contact the Patent Office.

METHOD

The present invention relates to a prognostic method and prognostic means based on polymorphisms in the vitamin D receptor gene. In particular, the present invention 5 relates to a method for determining susceptibility to heart disease by screening for polymorphisms in the vitamin D receptor gene.

Heart disease such as myocardial infarction is a complex phenotype caused by interaction of a number of genetic and environmental factors.¹ Advances in molecular 10 genetics have led to the identification of a number of genetic risk factors for heart disease, such as gene variants involved in thrombosis^{2,3} and lipid metabolism⁴. However, another important risk factor for heart disease such as myocardial infarction 15 is a deregulated calcium homeostasis which is required for maintenance of neuromuscular activity⁵. Disturbances of the calcium balance have been implicated in hypertension, ventricular hypertrophy, aortic calcification and arrhythmias all of which are important risk factors associated with myocardial infarction.

Vitamin D is a potent regulator of bone and calcium homeostasis, as well as of cellular 20 differentiation and replication in many tissues, and mediates its effects through the vitamin D receptor (VDR). Cloning of the vitamin D receptor has shown it to be a member of the ligand-activated superfamily, which are natural regulators of a number 25 of physiological and developmental processes. The major physiological function of the active metabolite of vitamin D, i.e. 1,25(OH)₂D₃, is to maintain serum calcium levels in the normal range. This is accomplished by stimulating the small intestine to increase its efficiency of absorbing calcium from the diet and to mobilize calcium stores from bone. The action of the hormone is mediated by the vitamin D receptor (VDR), a steroid transcription factor that mediates hormone response by initiating gene transcription and mRNA translation⁶. Involvement of the vitamin D endocrine system in the etiology of heart disease was suggested by epidemiological studies

which found serum levels of 25-hydroxyvitamin D₃ to be reduced in myocardial infarction patients compared with control individuals⁷. Furthermore, congestive heart failure associated with vitamin D deficiency has been described in a three and a half month old child⁸. The involvement of the vitamin D endocrine system in cardiac arrhythmias was suggested by case reports involving the observation of atrial flutter in a foetus with X-linked vitamin D resistant rickets⁹ and sick sinus syndrome that was cured by administration of vitamin D to a 77-year old caucasian woman¹⁰. Further studies in the rat have also demonstrated an important role of the vitamin D endocrine system in cardiovascular function, wherein vitamin D₃-deficiency can lead to large increases in the contractile function of the heart¹⁷⁻²⁰.

The above studies establish a link between the vitamin D endocrine system and risk of heart disease in an individual. However, the genetic component of heart disease such as myocardial infarction or cardiac arrhythmia is poorly defined. As a result, methods of diagnosing risk of heart disease, or treating those suffering from heart disease, are restricted.

Thus, in a first aspect of the present invention, there is provided a method of determining susceptibility to heart disease in a subject, said method comprising analysing genetic material of a subject to determine which allele(s) of the vitamin D receptor is/are present.

The vitamin D receptor gene (12q12) comprises inherited polymorphisms between exon 7 and the 3' UTR of the VDR gene, as shown in Figure 1. These alleles are denoted B/b, A/a and T/t for restriction enzyme sites *Bsm*I, *Apa*I and *Taq*I respectively (or enzymatic or chemical procedures with similar specificities), where a lower case letter denotes the presence of a wild type restriction site which is capable of being cleaved, and a capital letter denotes the presence of a mutant restriction enzyme site which is not capable of being cleaved by the relevant restriction enzyme. For the

purposes of the present invention, determination of which alleles are present in a particular gene may be referred to as determining the genotype of a subject for a particular gene. It is apparent from the above that each copy of the vitamin D receptor gene will comprise a specific combination of the three alleles, this combination being referred to as the haplotype of the gene. For example, the haplotype may be baT, indicating the presence of cleavable *BsmI* and *Apal* sites, and a non-cleavable *TaqI* site. Direct haplotyping of the VDR gene has allowed five different haplotypes to be determined, of which three are common.¹⁶

The present invention is based on the discovery of a genetic component of heart disease. It has now been shown that presence of the b, a or T alleles, and in particular the baT haplotype of the vitamin D receptor is/are associated with increased risk of heart disease. Thus, the presence of such alleles of the vitamin D receptor gene may be used to determine susceptibility to heart disease.

Preferably, the method of the first aspect of the present invention comprises the additional step of determining whether the alleles present are associated with risk of heart disease. This may be performed by comparing the alleles present in the genetic material of the subject with genotypes of the vitamin D receptor having known degrees of risk of heart disease. For example, a visual aid detailing alleles and their relative risk of heart disease may be used to determine whether the genotype of the subject is associated with a high or low risk of heart disease.

The first aspect of the present invention may also comprise the additional step of determining aspects of calcium metabolism, such as calcium levels, in a subject. Preferably, the daily calcium intake is measured. This feature of the first aspect is based on the observation that the correlation between vitamin D receptor genotype and heart disease may be dependent upon dietary calcium intake.

The method of the first aspect may be performed *in vitro*. Preferably, the method is performed on tissue or fluid removed from the body of the subject. Thus, the present invention relates to a non-invasive method, the results of which provide an indication of susceptibility to heart disease but do not lead to a diagnosis upon which an immediate medical decision regarding treatment has to be made.

5

The present invention may be performed on any subject for whom it is desirable to determine risk of heart disease. Preferably, the subject may be a mammal. Most preferably, the method of the first aspect is performed on humans, preferably male.

10

Heart disease may include atrial or ventricular hypertrophy, aortic calcification, arrhythmia, hypertension, and most importantly, myocardial infarction.

15

In a preferred feature of the first aspect of the present invention, there is provided a method of determining susceptibility to heart disease, said method comprising analysing genetic material of a subject to determine which of the B/b, A/a or T/t alleles of the vitamin D receptor gene is/are present. The method may comprise determining whether one or more of the above alleles is present. The subject may further be classified as being heterozygous or homozygous for one or more alleles. Preferably, the method comprises the additional step of determining whether the allele(s) present is/are associated with risk of heart disease, wherein presence of the b, a or T alleles is associated with increased risk of heart disease. Homozygosity for the a, b or T allele may further increase the susceptibility to heart disease in a subject.

20

25

30

In a preferred feature of the first aspect, there is provided a method of determining susceptibility to heart disease in a subject, said method comprising analysing the genetic material of a subject to determine the haplotype of the *BsmI*, *ApaI* and *TaqI* alleles at the vitamin D receptor. Preferably, said method comprises determining whether the haplotype of the subject is associated with risk of heart disease, wherein the haplotype baT is associated with high risk of heart disease. A subject homozygous

for said haplotype may be at a higher risk of heart disease than those heterozygous for the haplotype.

In a preferred feature of the first aspect, there is provided a method of determining 5 susceptibility to heart disease, said method comprising the additional step of determining the copy number of the B/b, A/a or T/t alleles or haplotype of the vitamin D receptor, where an increase in copy number is associated with increased risk of heart disease.

10 The present invention may be performed using any suitable method known in the art. Preferably, a tissue or fluid sample is first removed from a subject. Examples of suitable samples include blood, mouth or cheek cells, and hair samples containing roots. Other suitable samples would be known to the person skilled in the art. The 15 genetic material is then extracted from the sample for diagnosis, using any suitable method. The genetic material may be DNA or RNA, although preferably DNA is used. For example, the DNA may be extracted using the technique described in Sambrook *et al* (Molecular Cloning- A Laboratory Manual, Cold Spring Harbor Laboratory Press). Determination of the genotype of a subject may then be carried out using the extracted DNA, employing any one of the following techniques:

20

- Southern blot analysis following digestion with one or more appropriate restriction enzymes.
- PCR amplification followed by digestion with one or more appropriate restriction enzymes and, optionally, separation of digestion products by gel electrophoresis.
- 25 ▪ Sequencing of a relevant gene fragment by any suitable method.
- Visualization of heteroduplex patterns, for example on PAA or agarose gels, where different patterns may indicate the presence of one or more specific alleles.
- Separation of DNA fragments using denaturing gradient gels, wherein the degree of separation will depend upon the presence or absence of one or more polymorphic restriction sites.

30

- Separation using SSCP analysis, the patterns of which will depend upon the presence or absence of one or more polymorphic restriction sites.
- Use of allele specific oligonucleotides, hybridization patterns of which will be specific for various combinations of alleles.
- 5 ▪ Methods such as OLA, Taqman or dot-blot for the detection of known mutations.
- Visualization of DNA sites using fluorescent labelled probes for alleles of interest.
- RFLP analysis

Where it is desirable to use particular restriction enzymes in performing the present
10 invention, the skilled person will understand that enzymatic or chemical procedures having similar specificities may also be used. For example, restriction enzymes having similar specificity (isoschizomers) to those described herein may be used, or chemical degradation procedures with DNA or RNA cutting specificity.

15 Other techniques suitable for determining the genotype of a subject may be used in the present invention.

Where the haplotype of a gene is to be determined, it is preferable to use a direct
20 haplotyping method, as described in Uitterlinden *et al*¹⁶. In such a method, the relevant portion of the gene is amplified and then subjected to restriction enzyme digestion, in order to determine the presence or absence of restriction enzyme sites. Thus, for example, where the haplotype of the vitamin D receptor gene is to be determined, the portion of the gene between exon 7 and the 3' UTR may be amplified, and the amplified DNA digested with the *BsmI*, *ApaI* or *TaqI* restriction enzymes. Gel
25 analysis may then be used to determine which alleles are present.

30 Preferably, a fragment may be amplified using polymerase chain reaction (PCR) techniques, to produce copies which, where the fragment is of the vitamin D receptor, are at least 1000 base pairs in length, and most preferably at least 1800 base pairs in length. PCR techniques are well known in the art, and it is within the ambit of the

skilled person to identify primers for amplification of the appropriate region of the above genes, namely the region from exon 7 to the 3' UTR of the vitamin D receptor gene. PCR techniques are described in EP-A-0200362 and EP-A-0201184.

5 In a preferred feature of the first aspect, there is provided a method of determining susceptibility to heart disease in a subject, said method comprising amplifying a fragment comprising a portion of the region from exon 7 to the 3' UTR of the vitamin D receptor gene, and determining which allele(s) in the vitamin D receptor is/are present. Primers suitable for amplification of said portion of the vitamin D receptor 10 gene would be readily available to a person skilled in the art. Examples of such primers include:

1. 5'-CAACCAAGACTACAAGTACCGCGTCAGTGA-3' and/or
2. 5'-GCAACTCCTCATGGCTGAGGTCTC-3'

15 Where the amplified portion of the gene is larger than the above defined portion of the vitamin D receptor gene containing the relevant alleles, it is preferable to avoid the inclusion of gene sequences which comprise any one of the *BsmI*, *Apal* or *TaqI* restriction sites.

20 In a second aspect of the present invention, there is provided a method of therapy, said method comprising treating a subject diagnosed as being at risk of heart disease, to reduce the risk of heart disease. Preferably, the subject is diagnosed as being at risk of heart disease in accordance with the first aspect of the present invention.

25 Therapy may in the form of preventative or palliative care. Suitable treatments include modifications to lifestyle, regular exercise and changes in diet. Suitable treatments, including pharmaceutical preparations, would be known to physicians and persons skilled in the art. Examples include ACE inhibitors, beta-blockers, calcium or 30 vitamin D preparations, magnesium sulphate, thrombolytics and analgeasics.

In a third aspect of the present invention, there is provided a method of predicting the response of a subject to treatment, said method comprising analysing genetic material of a subject to determine which allele(s) of the vitamin D receptor gene is/are present.

5 Preferably, the method includes first determining whether the subject is susceptible to heart disease. Where a subject has been determined as susceptible to heart disease, the method may further comprise administering the appropriate treatment. The present aspect of the invention is based on the observation that agents, such as calcium channel blockers, which are useful for treatment of a variety of cardiovascular diseases

10 may also be associated with cardiovascular morbidity in some cases. The effect of an agent may therefore depend on the underlying cause of the heart disease. For example, the presence of the b, a or T alleles of the vitamin D receptor may result in modification of calcium uptake, leading to impaired cardiovascular function. Thus, in such a case it would be preferable to avoid the use of calcium channel blocking agents.

15 In a fourth aspect of the present invention, there is provided use of a kit to determine which allele(s) of the vitamin D receptor gene is/are present, said kit comprising (i) one or more nucleic acid primer molecules for amplification of a portion of the vitamin D receptor gene, and (ii) means for determining which allele(s) is/are present

20 in said gene.

Preferably, the primer molecules are suitable for amplification of at least a portion of the region between exon 7 and the 3'UTR of the vitamin D receptor gene. Examples of suitable primers are described above.

25 Means for determining which allele(s) is/are present in the vitamin D receptor gene, may include any reagents or molecules necessary for use in any of the methods described above. For example, where PCR followed by DNA digestion is used, said means preferably include PCR reagents and one or more of the *BsmI*, *Apal*, or *TaqI*

30 restriction enzymes. Where the method employs Southern Blotting, heteroduplex

visualization, or fluorescent labelling techniques for example, probes which bind to the appropriate regions of the vitamin D receptor gene may be included. Where necessary, such probes may be labelled to allow detection, for example by nick-translation, radio- or fluorescent-labelling, or random primer extension whereby the non-labelled nucleotides serve as a template for the synthesis of labelled molecules.

5 Other methods of labelling probes are well known in the art.

In a preferred feature of the fourth aspect of the present invention, there is provided use of a kit further comprising means for indicating correlation between the genotype 10 of a subject and risk of heart disease. Said means may be in the form of a chart or visual aid, which indicate that presence of the b, a or T alleles or baT haplotype of the vitamin D receptor is/are associated with heart disease such as cardiac arrhythmia or myocardial infarction.

15 In a fifth aspect of the present invention, there is provided a kit for determining risk of heart disease in a subject, said kit comprising (i) one or more nucleic acid primer molecules for amplification of a portion of the vitamin D receptor gene; (ii) means for determining which allele(s) is/are present in said gene; and (iii) means for indicating correlation between the allele(s) and risk of heart disease.

20 In a preferred feature of the fifth aspect, the kit may also comprise DNA control samples, for comparison with DNA sequences of a subject. The control samples may comprise the sequence of one or more alleles of the vitamin D receptor gene, or may comprise the sequence of various haplotypes.

25 Preferred features of each aspect of the present invention are as for each other aspect, mutatis mutandis.

30 The present invention will now be described in detail with reference to the following examples and figures in which:

FIGURE 1 is a schematic presentation of the region between exon 7 and the 3' UTR of the vitamin D receptor gene.

5 **Example 1 - Relationship between alleles of the vitamin D receptor gene and myocardial infarction, cardiac arrhythmias and dietary calcium intake in the elderly**

METHODS

10 Study Subjects

The Rotterdam Study is a population-based cohort study of 7983 subjects aged 55 or more years, residing in the Ommoord district of the city of Rotterdam in the Netherlands. The study was designed to document the occurrence of disease in the elderly in relation to several potential determinants.¹⁵ A total of 10,275 persons, of whom 9161 (89 percent) were living independently, were invited to participate in the 15 study in 1991. In the independently living subjects, the overall response rate was 77 percent for home interview and 71 percent for examination in a research centre, including measurement of anthropometric characteristics and blood sampling. The Rotterdam Study was approved by the Medical Ethics Committee of the Erasmus University Medical School and written informed consent was obtained from each 20 subject.

25 The analysis of the association between *VDR* genotype, myocardial infarction and arrhythmias was performed in a subgroup of men and women participating in the study. Baseline measurements, collected from 1990-1993, included an electrocardiogram (ECG), history of cardiovascular disease, established cardiovascular risk factors, and use of medication. A digitally stored ECG was available for 5931 independently living 30 subjects from the study, but 1453 of these were excluded on the basis of age (>80 yrs), use of a walking aid, diabetes mellitus or use of diuretic, estrogen, thyroid hormone or cytostatic drug therapy. From the 4478 remaining subjects, we studied a random

sample of 2000 men and women aged 55 to 80 years. ECG data or DNA samples were not available for 22 subjects, resulting in a final study group of 1978 subjects.

Measurements

5 Height and weight were measured at the initial examination in a standing position without shoes. Myocardial infarction, including so called silent myocardial infarctions, was confirmed by ECG analysis. For 1725 men and women (87%) data was available on dietary intakes of calcium (mg/day) during the preceding year. These were assessed by food frequency questionnaire and adjusted for energy intake. Age at menopause
10 and current cigarette smoking were assessed by questionnaire. Cardiac arrhythmias were diagnosed by ECG analysis and included the occurrence of sinus irregularities, atrial flutter, and atrial fibrillation and were recorded, confirmed and classified by a physician. For 976 subjects data on cardiac arrhythmias were available (49%). Of these subjects, data on dietary calcium intake were available for 899 subjects (92%).

15

Determination of VDR Genotypes

Genomic DNA was extracted from peripheral venous blood samples according to standard procedures and the anonymous polymorphisms were detected by PCR as previously described¹⁶. Three anonymous polymorphic restriction enzyme recognition sites at the 3' end of the VDR gene, i.e. for BsmI, ApaI, and TaqI, were assessed in relation to each other by a direct molecular haplotyping PCR procedure which we developed¹⁶. This allows us to determine ophase of the alleles at each of the RFLP loci and as a result three frequent haplotype alleles are discerned, encoded 1 (baT; frequency 48%), 2 (BAT; frequency 40%), 3 (bAT; frequency 10%) combining to six genotypes encoded 11, 12, 13, 22, 23, and 33. Detailed information on haplotype alleles and genotype frequencies in the Rotterdam Study can be found elsewhere¹⁶. The PCR reaction mixture of 25 microlitres contained 100 ng of genomic DNA, 50 mM KCl, 10 mM Tris-HCl (pH 8.3), 1.5 mM MgCl₂, 0.2 mM dideoxynucleotidetriphosphates, 150 ng of each primer, and 0.2 unit of Super Taq

polymerase (HT Biotechnology, Cambridge, UK). The reactions were performed in a DNA thermocycler (model 480, Perkin Elmer) with a cycling protocol of 94°C, 60°C, and 72°C for 1 min each, for 28 cycles. Ten microlitres of PCR product were simultaneously digested with 5 units BsamI, 5 units ApaI, and 5 units TaqI restriction enzyme and 2 microlitres of a 10x concentrated buffer (containing 150 mM Tris-HCl, pH 7.5, 250 mM NaCl, and 35 mM MgCl₂) by incubating for 30 min at 37°C, followed by a 30 min incubation at 65°C. The digestion products were analysed on a 1.2 percent agarose gel run in 44.5 mM Tris, 44.5 mM boric acid, 1 mM Na₂EDTA) for 300 volt-hours. Separation patterns were documented by Polaroid photography under UV-illumination (302 nm). To confirm the accuracy of genotyping, repeat analysis was performed on 200 random samples. No discrepancies were found.

Statistical Analysis

Relevant clinical variables were compared between the genotype groups by analysis of covariance and multiple linear regression was used to adjust values for confounding factors such as age, anthropometric variables, smoking. For the comparisons we made reference, heterozygote and homozygote groups for each of the VDR alleles. For example, for VDR allele1 the groups comprised 22, 23, and 33 for the reference group, 12, and 13 for the heterozygote group, and 11 for the homozygote group. The Chi-squared test was used to test for genotype distribution in men and women with and without myocardial infarction and with and without cardiac arrhythmias. Odds ratios (with 95 percent confidence interval) were calculated by multivariate logistic regression analysis to estimate the relative risk of myocardial infarction and cardiac arrhythmias. A multivariate regression model was designed to test for the influence of dietary calcium intake on the relationship between VDR genotype and myocardial infarction and VDR genotype with cardiac arrhythmias. For this analysis, subjects were categorised into groups of calcium intake defined by quartiles of the population. An interaction term was included in the regression analysis defined as gene allele dose multiplied with calcium-intake. P values for all statistical tests were two-sided.

RESULTS

Characteristics of the Study Subjects

Anthropometric, and dietary measurements in the men and women with the six VDR genotypes are shown in Table 1. The allele frequencies (1=48.6%; 2=40.6%; and 3=10.8%) and the distribution of genotypes were similar to that reported in a previous study²². There was no difference with regard to age, dietary calcium intake, smoking habits, body mass index, serum HDL-cholesterol , and serum cholesterol in the six genotype groups.

10

Association with Myocardial Infarction

The distribution of myocardial infarction by VDR genotype in men and women is shown in Table 2. There was modest over-representation of both men and women carrying the VDR haplotype allele 1 as compared with subjects without the allele although this did not reach significance. Logistic regression analysis showed that subjects in the heterozygous group had a 1.2 fold increased risk for myocardial infarction and the subjects in the homozygous group had a 1.5-fold increased risk, as compared with subjects not carrying the VDR allele 1. The gene dose effect was 1.2-fold increased risk per copy of the VDR allele 1. The gene dose effect was larger for women when compared to men (RRs were 1.1 (95% confidence interval 0.9 – 1.4) for men and 1.5 (95% confidence interval 1.1 – 2.2) for women) but the interaction term of sex times genotype in the regression model was not significant ($P=0.65$). The relative risks for myocardial infarction did not change after adjustment for potential confounding factors such as age, body mass index and smoking in the regression analysis. For VDR haplotype allele 2 and haplotype allele 3 no significant differences were observed (data not shown).

Modification of the Association by Dietary Calcium Intake

When the study subjects were stratified according to quartiles of dietary calcium intake, the differences in distribution of myocardial infarction by VDR genotype were restricted to individuals with the highest intake of calcium (1302 mg/day or more).

5 These differences corresponded with 5.4-fold increased risk for heterozygote carriers and 8.3 -fold increased risk for homozygote carriers of VDR haplotype allele 1. Considering age, genotype and myocardial infarction together in a multivariate regression model, we found a significant modifying effect of dietary calcium intake on the VDR genotype effect on risk for myocardial infarction ($P=0.04$).

10

Association with Cardiac Arrhythmias

We subsequently analysed the distribution of cardiac arrhythmias by VDR genotype. We did not observe differences in the distribution of arrythmias by VDR genotype in the total study group. However, in the group with high calcium intake (1302 mg/day or more) there was significant over-representation of cardiac arrythmias for subjects carrying the VDR haplotype allele 1 as compared with subjects without the allele 1 (Table 4).

15 Logistic regression analysis showed that subjects in the heterozygous group had a 1.6 fold increased risk for cardiac arrhythmias and the subjects in the homozygous group had a 3.6-fold increased risk, as compared with subjects not carrying the VDR allele 1. The relative risks for cardiac arrythmia did not change after adjustment for potential confounding factors such as age, body mass index and smoking in the regression analysis. Also when myocardial infarction was entered into the model the relative risks 25 did not change.

DISCUSSION

30 Polymorphisms of the vitamin D receptor gene are associated with the risk of myocardial infarction. The increased risk was especially notable in subjects with a

high dietary calcium intake. In this group, we also observed VDR haplotype 1 to be associated with increased risk of cardiac arrhythmias. Cardiac arrhythmias have been frequently observed following a myocardial infarct event. However, the VDR genotype dependent risk for cardiac arrhythmias we observe, is independent of myocardial infarction. This indicates that the genotype effect we observe for myocardial infarction is not mediated by the genotype effect on cardiac arrhythmias.

Our results indicate a genetic susceptibility of the risk for myocardial infarction and cardiac arrhythmias in subjects with a relatively high dietary calcium intake. This strongly suggest that the VDR genotype effect on myocardial infarction and cardiac arrhythmias is mediated by an effect on calcium transport. Treatment with calcium and vitamin D has been indicated for arrhythmias and hypertrophy and has been shown to improve myocardial function.

15 The hormone vitamin D has been shown to influence the Ca-flux in a variety of cell types including heart muscle cells. These rapid effects of the hormone are suggested to involve a non-genomic pathway by which the plasmalemmal VDR is coupled to plasma membrane Ca²⁺ channel proteins.

20 Calcium channel blocking agents are indicated for the treatment of a variety of cardiovascular diseases, including cardiac arrhythmias, but have also been found associated with increased cardiovascular morbidity^{21,22}. Therefore, the modification by calcium intake of the VDR genotype dependent effect on myocardial infarction we observe opens the possibility that genotyping at the VDR gene locus might be of value in predicting the response to treatment with some of these drugs. Prospective clinical trials are needed to investigate the clinical and therapeutic implications of our results.

25

REFERENCES

1. Marian AJ. Genetic risk factors for myocardial infarction. *Curr Opin Cardiol* 1998;13:171-178.
- 5 2. Weiss EJ, Bray PF, Tayback M, et al. A polymorphism of a platelet glycoprotein receptor as an inherited risk factor for coronary thrombosis. *New Engl J Med* 1996;334:1090-1096.
- 10 3. Iacoviello L, Di Castelnuovo A, de Knijff P, et al. Polymorphisms in the coagulation factor VII gene and the risk of myocardial infarction. *New Engl. J Med* 1998;338:79-85.
4. Kuivenhoven JA, Jukema W, Zwinderman AH, et al. The role of a common variant of the cholesteryl ester transfer protein gene in the progression of coronary atherosclerosis. *New Engl. J Med* 1998;338:86-93.
- 15 5. Lind L, Skarfors E, Berglund L, Lithell H, and Ljunghall. Serum calcium: a new, independent, prospective risk factor for myocardial infarction in middle-aged men followed for 18 years. *J Clin Epidemiol* 1997;50:967-973.
6. Haussler MR, Haussler CA, Jurutka PW, et al. The vitamin D hormone and its nuclear receptor: molecular actions and disease states. *J Endocrinol* 1997;154:S57-S73.
- 20 7. Scragg R, Jackson R, Holdaway IM, Lim T, and Beaglehole R. Myocardial infarction is inversely associated with plasma 25-hydroxyvitamin D₃ levels: a community based study. *Int J Epidemiol* 1990;19:559-563.
8. Brunvand L, Haga P, Tangsrud SE, and Haug E. Congestive heart failure caused by vitamin D deficiency? *Acta Paediatr* 1995;84:106-108.
- 25 9. Vintzileos AM, Campbell WA, Soberman SM, and Nochimson DJ. Fetal atrial flutter and X-linked dominant vitamin D resistant rickets. *Obstet Gynecol*. 1985;65:39S-44S.
10. Kessel L. Sick sinus syndrome cured by ... vitamin D? *Geriatrics* 1990;45:83-85.

11. Cooper GS, Umbach DM. Are vitamin D receptor polymorphisms associated with bone density ? J Bone Miner Res 1996;11:1841-9.
12. Carling T, Kindmark A, Helleman P, et al. Vitamin D receptor genotypes in primary hyperparathyroidism. Nature Med 1995;1:1309-1311.
- 5 13. Taylor JA, Hirvonen A, Watson M, Pittman G, Mohler JL, and Bell DA. Association of prostate cancer with vitamin D receptor gene polymorphism. Canc. Res. 1996;56:4108-4110.
14. Uitterlinden AG, Burger H, Huang Q, et al. Vitamin D receptor genotype is associated with radiographic osteoarthritis at the knee. J Clin Invest 1997;100:259-10 263.
15. Hofman A, Grobbee DE, de Jong, PTVM, van den Ouwehand FA. Determinants of disease and disability in the elderly: the Rotterdam Elderly Study. Eur J Epidemiol 1991;7:403-22.
16. Uitterlinden AG, Pols HAP, Burger H, et al. A large scale population based study of the association of vitamin D receptor gene polymorphisms with bone mineral density. J Bone Miner Res 1996;11:1242-1248.
17. O'Connell TD, Weishaar RE, and Simpson RU. Regulation of myosin isozyme expression by vitamin D3 deficiency and 1,25-dihydroxyvitamin D3 in the rat heart. Endocrinology 1994;134: 899-905
- 20 18. Weishaar RE, Kim SN, Saunders D, and Simpson RU. Involvement of vitamin D3 with cardiovascular function III. Effects on physical and morphological properties. Am J Physiol 1990;258 (Endocrinol. Metab. 21):E134-E142.
19. Weishaar RE, and Simpson RU. Involvement of vitamin D3 with cardiovascular function II. Direct and indirect effects. Am J Physiol 1987;253 (Endocrinol. Metab. 16):E675-E683.
- 25 20. Weishaar RE, and Simpson RU. Vitamin D3 and cardiovascular function in rats. J Clin Invest 1987;79:1706-1712.
21. Psaty BM, Heckbert SR, Koepsell TD, et al. The risk of myocardial infarction associated with antihypertensive drug therapies. JAMA 1995; 271:620-625.
- 30 22. Michalewicz L, and Messerli FH. Cardiac effects of calcium antagonists in systemic hypertension. Am J Cardiol. 1997; 79:39-46.

CLAIMS

1. A method of determining susceptibility to heart disease in a subject, said method comprising analysing genetic material of a subject to determine which allele(s) of the vitamin D receptor gene is/are present.
5
2. A method of determining susceptibility to heart disease according to claim 1, said method comprising analysing the genetic material of a subject to determine which of the B/b, A/a or T/t alleles of the *BsmI*, *ApaI* or *TaqI* sites of the vitamin D receptor
10 are present.
3. A method of determining susceptibility to heart disease according to claim 1, said method comprising analysing the genetic material of a subject to determine the haplotype of the *BsmI*, *ApaI* or *TaqI* alleles of the vitamin D receptor.
15
4. A method according to claim 3 wherein the haplotype may be determined by amplification of a relevant portion of the vitamin D receptor gene, followed by restriction enzyme digestion.
- 20 5. A method of determining susceptibility to heart disease according to claims 1 to 4, said method comprising determining the copy number of the B/b, A/a or T/t alleles or haplotype of the vitamin D receptor.
- 25 6. A method according to any one of the preceding claims further comprising determining whether the allele(s) present is/are associated with risk of heart disease.
7. A method according to claim 6 comprising comparing the allele(s) present in the genetic material of the subject with those known to be associated allele(s) of vitamin D receptor genotypes having known degrees of risk of heart disease

8. A method according to any one of the previous claims wherein said method further comprises determining aspects of calcium metabolism in a subject.

9. A method according to claim 8, wherein daily calcium intake of a subject is measured.

10. A method according to any one of the preceding claims, wherein said method is performed *in vitro*.

10 11. A method according to claim 10 wherein said method is performed on blood or tissue samples of a subject.

12. A method according to any one of the preceding claims wherein the subject is a mammal.

15

13. A method according to claim 12 wherein the subject is a human.

14. A method according to claims 12 or 13 wherein the subject is male.

20 15. A method according to any one of the previous claims for determining susceptibility of a subject to atrial or ventricular hypertrophy, aortic calcification, arrhythmia, myocardial infarction, or hypertension.

25 16. A method according to any one of the preceding claims further comprising treating the subject to reduce the risk of heart disease.

17. A method according to claim 16 wherein suitable treatments may include modifications to lifestyle, regular exercise, changes in diet or pharmaceutical preparations.

30

18. A method of predicting the response of a subject to treatment, said method

comprising analysing genetic material of a subject to determine which allele(s) of the vitamin D receptor gene is/are present.

19. A method according to claim 18 wherein said subject is first diagnosed as
5 being susceptible to heart disease in accordance with any one of claims 1 to 17.

20. A method according to claims 18 or 19 further comprising administering the appropriate treatment.

10 21. Use of a kit to determine susceptibility to heart disease in a subject, said kit comprising (i) one or more nucleic acid primer molecules for amplification of a portion of the vitamin D receptor gene, and (ii) means for determining which allele(s) of said gene is/are present.

15 22. A kit for determining susceptibility to heart disease in a subject, said kit comprising (i) one or more nucleic acid primer molecules for amplification of a portion of the vitamin D receptor gene; (ii) means for determining which allele(s) of said gene is/are present; and (iii) means for indicating correlation between said allele(s) and risk of heart disease.

20

23. A kit according to claim 22, said kit comprising DNA control samples, for comparison with DNA sequences of a subject.

Table 1. Characteristics of the Population According to *VDR* Genotype

CHARACTERISTIC†	<i>VDR</i> GENOTYPE				P-VALUE
	11	12	13	22	
Number (%)	493 (24.9)	735 (37.2)	202 (10.2)	351 (17.7)	170 (8.6) 27 (1.4) 0.93§
Age (years)	67.0 ± 6.8	67.1 ± 6.8	67.2 ± 7.1	67.0 ± 7.1	66.9 ± 6.8 67.0 ± 7.1 0.78¶
Body Mass Index (kg/m ²)	26.1 ± 3.7	26.0 ± 3.3	25.8 ± 3.6	26.1 ± 3.4	26.0 ± 3.2 25.5 ± 2.9 0.78¶
Dietary calcium-intake (mg/day)	1116 ± 350	1122 ± 364	1122 ± 356	1092 ± 369	1094 ± 342 1158 ± 254 0.78¶
Serum HDL-cholesterol (mmol/l)	1.34 ± 0.37	1.35 ± 0.36	1.36 ± 0.34	1.35 ± 0.37	1.32 ± 0.33 1.36 ± 0.38 0.91
Serum cholesterol (mmol/l)	6.68 ± 1.21	6.63 ± 1.26	6.64 ± 1.16	6.60 ± 1.19	6.59 ± 1.21 6.60 ± 0.96 0.95
Current Smokers (%)	130 (26.4)	172 (23.4)	45 (22.3)	78 (22.2)	40 (23.5) 6 (22.2) 0.83¶

† Values are means ± standard deviation; BMI is weight divided by the square height

§ P-value for ANOVA

¶ P-value for Chi-2 test

Table 2. Myocardial Infarction According to *VDR allele 1* Genotype

	Men		Women		All	
	MI (%)	Total	MI (%)	Total	MI (%)	Total
Total	151 (15.8)	954	62 (6.1)	1024	213 (10.8)	1978
by <i>VDR allele 1</i> genotype						
<i>Reference†</i>	39 (14.7)	266	10 (3.5)	282	49 (8.9)	548
<i>Heterozygotes</i>	69 (15.4)	449	31 (6.4)	488	100 (10.7)	937
<i>Homozygotes</i>	43 (18.0)	239	21 (8.3)	254	64 (13.0)	493
χ^2	1.18		5.38		4.43	
P-VALUE	0.55		0.07		0.11	
Odds Ratios for Myocardial Infarct by <i>VDR allele 1</i> genotype [95% CI]						
Crude						
<i>Reference</i>	1.00		1.00		1.00	
<i>Heterozygotes</i>	1.07 [0.72 - 1.71]		1.86 [0.90 - 3.85]		1.23 [0.86 - 1.76]	
<i>Homozygotes</i>	1.28 [0.80 - 2.05]		2.48 [1.15 - 5.39]		1.53 [1.03 - 2.27]	
per copy <i>VDR 1</i> allele	1.13 [0.89 - 1.44]		1.53 [1.07 - 2.20]		1.24 [1.02 - 1.51]	
Age-, BMI-adjusted						
<i>Reference</i>	1.00		1.00		1.00	
<i>Heterozygotes</i>	1.11 [0.72 - 1.71]		1.77 [0.85 - 3.68]		1.22 [0.85 - 1.75]	
<i>Homozygotes</i>	1.33 [0.82 - 2.14]		2.45 [1.12 - 5.34]		1.55 [1.04 - 2.30]	
per copy <i>VDR 1</i> allele	1.15 [0.91 - 1.47]		1.53 [1.06 - 2.22]		1.25 [1.02 - 1.52]	

† "Reference" includes VDR genotypes 22, 23, 33; "Heterozygotes" includes 12, 13; "Homozygotes" includes 11

Table 3. Myocardial Infarction According to *VDR allele 1* Genotype by Quartiles of Dietary Calcium Intake

	< 877 mg/day			> 877, < 1076			> 1076, < 1302			≥ 1302		
	MI (%)	Total	MI (%)	Total	MI (%)	Total	MI (%)	Total	MI (%)	Total	MI (%)	Total
Total	43 (10.0)	432	50 (11.6)	431	45 (10.4)	432	49 (11.4)	430				
by <i>VDR allele 1</i> genotype												
Reference†	13 (9.9)	131	14 (12.5)	112	12 (9.6)	125	3 (2.6)	114				
Heterozygotes	21 (10.5)	200	24 (11.8)	204	21 (10.0)	210	26 (12.6)	207				
Homozygotes	9 (8.9)	101	12 (10.4)	115	12 (12.4)	97	20 (18.3)	109				
χ^2	0.19		0.25		0.53		14.17					
P-VALUE	0.91		0.88		0.77		0.0008					

Odds Ratios for Myocardial Infarct by <i>VDR allele 1</i> genotype [95% CI]		
Crude		
Reference	1.00	1.00
Heterozygotes	1.09 [0.52 - 2.27]	0.93 [0.46 - 1.89]
Homozygotes	0.90 [0.37 - 2.20]	0.82 [0.36 - 1.87]

† "Reference" includes VDR genotypes 22, 23, 33; "Heterozygotes" includes 12, 13; "Homozygotes" includes 11

Table 4.

Myocardial Arrhythmias According to *VDR allele 1* Genotype by Quartiles of Dietary Calcium Intake

	< 877 mg/day			> 877, < 1076			> 1076, < 1302			≥ 1302		
	MA (%)	Total	MA (%)	Total	MA (%)	Total	MA (%)	Total	MA (%)	Total	MA (%)	Total
Total	37 (12.1)	307	27 (9.2)	292	17 (5.6)	302	31 (10.1)	306				
by <i>VDR allele 1</i> genotype												
Reference†	16 (17.0)	94	6 (8.8)	68	7 (7.7)	91	5 (5.7)	88				
Heterozygotes	14 (10.1)	138	14 (10.1)	138	6 (4.4)	135	12 (8.5)	141				
Homozygotes	7 (9.3)	75	7 (8.1)	86	4 (5.3)	76	14 (18.2)	77				
χ^2	3.19		0.27		1.11		7.80					
P-VALUE	0.20		0.87		0.58		0.02					

Odds Ratios for Myocardial arrhythmias by <i>VDR allele 1</i> genotype [95% CI]		
Crude		
Reference	1.00	1.00
Heterozygotes	0.57 [0.26 - 1.23]	1.13 [0.41 - 3.12]
Homozygotes	0.51 [0.20 - 1.32]	0.92 [0.29 - 2.92]

† "Reference" includes *VDR* genotypes 22, 23, 33; "Heterozygotes" includes 12, 13; "Homozygotes" includes 11

